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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/844,935	04/27/2001	Alexander Munishkin	Q01/08C	4219

7590

12/23/2002

Attention: Anthony J. Janiuk, Esq.  
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New York, NY 10032

EXAMINER
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CHAKRABARTI, ARUN K

ART UNIT	PAPER NUMBER
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1634

DATE MAILED: 12/23/2002

Please find below and/or attached an Office communication concerning this application or proceeding.

# Office Action Summary

Application No.  
**09/844,935**

Applicant(s)  
**Muniskin**

Examiner  
**Arun Chakrabarti**

Art Unit  
**1634**



-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

## Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136 (a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

## Status

- 1) ☒ Responsive to communication(s) filed on Dec 9, 2002.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11; 453 O.G. 213.

## Disposition of Claims

- 4) ☒ Claim(s) 1-9 and 11-13 is/are pending in the application.
- 4a) Of the above, claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 1-8, 11, and 12 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☒ Claims 9 and 13 are subject to restriction and/or election requirement.

## Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on \_\_\_\_\_ is: a) ☐ approved b) ☐ disapproved by the Examiner.  
If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

## Priority under 35 U.S.C. §§ 119 and 120

- 13) ☐ Acknowledgement is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).  
a) ☐ All b) ☐ Some\* c) ☐ None of:  
1. ☐ Certified copies of the priority documents have been received.  
2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.  
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).  
\*See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgement is made of a claim for domestic priority under 35 U.S.C. § 119(e).  
a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgement is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

## Attachment(s)

- 1) ☐ Notice of References Cited (PTO-892) 4) ☐ Interview Summary (PTO-413) Paper No(s). \_\_\_\_\_
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948) 5) ☐ Notice of Informal Patent Application (PTO-152)
- 3) ☐ Information Disclosure Statement(s) (PTO-1449) Paper No(s). \_\_\_\_\_ 6) ☒ Other: **Detailed Action**

Art Unit: 1634

## **DETAILED ACTION**

### ***Election/Restriction***

1. Applicant argued that claims 9 and 13 of Groups II and III should be rejoined with claims 1-8, 11, and 12 of Group I. The traversal is on the ground(s) that there is no burden in examining the claims of Groups II, and III. This is not found persuasive because as the restriction makes clear, additional search of Groups II, and III would require review not only of the 1166 patents in class 536, subclass 22.1 for Group I, but also the 3801 patents in class 435, subclass 91.2 for Group II, and the 9976 patents in class 435, subclass 6 for Group III. Review of these additional searches is prima facie evidence of burden which is not rebutted.

The requirement is still deemed proper and is therefore made FINAL.

### ***Claim Rejections - 35 USC § 102***

2. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

3. Claims 1-5, and 7 are rejected under 35 U.S.C. 102(b) as being anticipated by Marsh et al.(Nucleic Acids Research, (1988), 16 (3), pages 981-995).

Art Unit: 1634

Marsh et al. teaches a method of determining the presence or absence of a target molecule (abstract) comprising the steps of:

a) providing a first RNA molecule which can bind to a target molecule and has the formula:

5'-A-B-C-D-E-3'; (Figure 1, 2 and 4)

wherein A is a section of the RNA molecule having 10-10,000 nucleotides which section is, with another sequence, E, replicated by an RNA replicase, the letter "B" denotes a section of the RNA molecule having approximately 1 to 50000 nucleotides which section, with another sequence D, binds the target molecule under binding conditions, the letter "C" denotes a section of the RNA molecule having approximately 1 to 10000 nucleotides, the letter "D" denotes a section of the RNA molecule having approximately 1 to 50000 nucleotides which section, with another B, binds the target molecule under binding conditions, the section B and D, in combination, comprise in total at least 10 nucleotides, the first RNA molecule, with sections B and D bound to target, is acted upon by the RNA replicase to form a second RNA molecule, said second RNA molecule has the following formula:

5'-E'-X-A'-3';

wherein, E' is the complement to E, and A' is the complement to A, and the letter "X" denotes the complement of parts of the sections B and D which may be replicated, or the letter denotes the direct bond between sections E' and A', and second RNA molecule is replicated by the RNA

Art Unit: 1634

replicase under replicating conditions and combining first RNA molecule with a sample (Figure 1, 2, 3 and 4 and Materials and Methods, page 983, lines 12-25);

b) imposing binding conditions on a sample potentially containing target molecules in the presence of first RNA molecule, in the presence of the target molecule, first RNA molecules forms a target-first RNA molecule complex to form a first modified sample (Figure 2, 3 and 4 );

c) imposing RNA replicase reaction conditions on the first modified sample, in the presence of an RNA replicase, to form second RNA molecule in the presence of target to make a second modified sample (Materials and Methods, page 983, lines 12-25);

d) monitoring second modified sample for the presence of the second RNA molecule or its complement, which presence or absence is indicative of the presence or absence of the target molecule (Materials and Methods, page 983, lines 25-32 and Table 1).

Marsh et al. teaches that section "C" may serve as a non base-paired spacer to facilitate access of the replicase to the promoter (page 990, lines 9-10).

### ***Claim Rejections - 35 USC § 103***

4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Art Unit: 1634

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CAR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103© and potential 35 U.S.C. 102(f) or (g) prior art under 35 U.S.C. 103(a).

5. Claims 6 and 8 are rejected under 35 U.S.C. 103(a) as being unpatentable over Marsh et al.(Nucleic Acids Research, (1988), 16 (3), pages 981-995) in view of Spiegelman (U.S. Patent 3,444,043) (May 13, 1969).

Marsh et al teaches the a composition of claims 1-7 as described above.

Marsh et al. does not teach a composition by providing paired RNA molecules.

Marsh et al does not teach section "C" of the RNA molecule which section is capable of preventing the replication of the first molecule by the RNA replicase (abstract and column 7, lines 8-44).

Spiegelman teaches the customized preparation of RNA templates as he states, "An RNA template of an in vitro replicating system may be formed in situ. If one were, for example, to introduce foreign bases or nucleotides (e.g., analogues of known bases or nucleotides) into the replicating system, a mutant may be formed which would be the biologically active template for

Art Unit: 1634

replication with those same bases or nucleotides, in such instances, one would be synthesizing mutants in vitro in a known way (Column 5, lines 1-8)".

Spiegelman teaches section "C" of the RNA molecule which section is capable preventing the replication of the first molecule by the RNA replicase (abstract and column 7, lines 8-44).

It would have been *prima facie* obvious to one having ordinary skill in the art at the time the invention was made to substitute RNA template model of Spiegelman as the identification of target molecule in the method of Marsh et al., since Spiegelman et al. states "There is good evidence that the replicase recognizes the particular sequence of nucleotides at the beginning and at the end of the biologically active viral RNA template during the course of the replication. It is inferred from this recognition pattern that the intermediate portion of the RNA template is not essential to the direction of or instruction found in the replication mechanism studied. This suggests that the recognition sequences of nucleotides present at the beginning and end of a biologically active RNA template molecule can be selectively bonded to otherwise non-biologically active or non-viral RNA to produce a synthesized biologically active RNA product. It is thought that the RNA forms a circle and these two recognition sequences of the molecule overlap each other to provide double-stranded regions: such overlapped regions could afford, therefore, identification of the RNA molecule in a single, rapid scanning process (Column 4, lines 59-75)". An ordinary practitioner would have been motivated to combine the model of custom made RNA template of Spiegelman into the method of Marsh et al. in order to achieve the

Art Unit: 1634

express advantages noted by Spiegelman of a method which can provide identification of the RNA molecule in a single, rapid scanning process.

6. Claim 11 is rejected under 35 U.S.C. 103(a) as being unpatentable over Marsh et al. (Nucleic Acids Research, (1988), 16 (3), pages 981-995) in view of Stratagene Catalog (1988, Page 39).

Marsh et al. teach the compositions of claims 1-5, and 7 as described above in detail.

Marsh et al. do not teach the motivation to combine all the reagents for detecting an analyte in a sample in the form of a kit.

Stratagene catalog teaches a motivation to combine reagents into kit format (page 39).

It would have been *prima facie* obvious to one having ordinary skill in the art at the time the invention was made to combine the compositions of claims 1-5, and 7 of Marsh et al. into a kit format as discussed by Stratagene catalog since the Stratagene catalog teaches a motivation for combining reagents of use in an assay into a kit, "Each kit provides two services: 1) a variety of different reagents have been assembled and pre-mixed specifically for a defined set of experiments. Thus one need not purchase gram quantities of 10 different reagents, each of which is needed in only microgram amounts, when beginning a series of experiments. When one considers all of the unused chemicals that typically accumulate in weighing rooms, desiccators, and freezers, one quickly realizes that it is actually far more expensive for a small number of users to prepare most buffer solutions from the basic reagents. Stratagene provides only the quantities you will actually need, premixed and tested. In actuality, the kit format saves money and

Art Unit: 1634

resources for everyone by dramatically reducing waste. 2) The other service provided in a kit is quality control (page 39, column 1).

7. Claim 12 is rejected under 35 U.S.C. 103(a) as being unpatentable over Marsh et al.(Nucleic Acids Research, (1988), 16 (3), pages 981-995) in view of Spiegelman (U.S. Patent 3,444,043) (May 13, 1969) further in view of Stratagene Catalog (1988, Page 39).

Marsh et al. in view of Spiegelman expressly teach the method claims and assay reagents of claims 8 as described above in detail.

Marsh et al. in view of Spiegelman do not teach the motivation to combine all the reagents for detecting an analyte in a sample in the form of a kit.

Stratagene catalog teaches a motivation to combine reagents into kit format (page 39).

It would have been *prima facie* obvious to one having ordinary skill in the art at the time the invention was made to combine the composition of Marsh et al. in view of Spiegelman into a kit format as discussed by Stratagene catalog since the Stratagene catalog teaches a motivation for combining reagents of use in an assay into a kit, "Each kit provides two services: 1) a variety of different reagents have been assembled and pre-mixed specifically for a defined set of experiments. Thus one need not purchase gram quantities of 10 different reagents, each of which is needed in only microgram amounts, when beginning a series of experiments. When one considers all of the unused chemicals that typically accumulate in weighing rooms, desiccators, and freezers, one quickly realizes that it is actually far more expensive for a small number of users to prepare most buffer solutions from the basic reagents. Stratagene provides only the quantities

Art Unit: 1634

you will actually need, premixed and tested. In actuality, the kit format saves money and resources for everyone by dramatically reducing waste. 2) The other service provided in a kit is quality control (page 39, column 1).

***Response to Amendment***

8. In response to amendment, restriction requirement as well as 102(b) and 103(a) rejections are hereby being properly maintained.

***Response to Arguments***

9. Applicant's arguments filed on December 9, 2002 have been fully considered but they are not persuasive.

In response to applicant's arguments against the references individually, one cannot show nonobviousness by attacking references individually where the rejections are based on combinations of references. See *In re Keller*, 642 F.2d 413, 208 USPQ 871 (CCPA 1981); *In re Merck & Co.*, 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986). Applicant argues that there is no motivation to combine the references. This argument is not persuasive, especially in the presence of strong motivation provided by Spiegelman et al. since Spiegelman et al. states "There is good evidence that the replicase recognizes the particular sequence of nucleotides at the beginning and at the end of the biologically active viral RNA template during the course of the replication. It is inferred from this recognition pattern that the intermediate portion of the RNA template is not

Art Unit: 1634

essential to the direction of or instruction found in the replication mechanism studied. This suggests that the recognition sequences of nucleotides present at the beginning and end of a biologically active RNA template molecule can be selectively bonded to otherwise non-biologically active or non-viral RNA to produce a synthesized biologically active RNA product. It is thought that the RNA forms a circle and these two recognition sequences of the molecule overlap each other to provide double-stranded regions: such overlapped regions could afford, therefore, identification of the RNA molecule in a single, rapid scanning process (Column 4, lines 59-75)". The same logic is applicable to other 103(a) references as well.

Applicant argues to withdraw the 102(b) rejections because Marsh does not teach some characteristic features of use of the composition of the instant invention. This argument is not persuasive. In response to applicant's argument that , a recitation of the intended use of the claimed invention must result in a structural difference between the claimed invention and the prior art in order to patentably distinguish the claimed invention from the prior art. If the prior art structure is capable of performing the intended use, then it meets the claim. In this case, Marsh clearly teaches that section "C" may serve as a non base-paired spacer to facilitate access of the replicase to the promoter (page 990, lines 9-10). In a claim drawn to a process of making, the intended use must result in a manipulative difference as compared to the prior art. See *In re Casey*, 152 USPQ 235 (CCPA 1967) and *In re Otto*, 136 USPQ 458, 459 (CCPA 1963).

Applicant argues to withdraw the 102(b) and 103(a) rejections because Marsh and Spiegelman do not teach some characteristic features of the instant invention. In response to

Art Unit: 1634

applicant's argument that the references fail to show certain features of applicant's invention, it is noted that the features upon which applicant relies (i.e., (I) the replicated strand is a different sequence (ii) applicant's claimed invention is not a mutation, and (iii) any specific type of single, rapid screening process ) are not recited in the rejected claim(s). Although the claims are interpreted in light of the specification, limitations from the specification are not read into the claims. See *In re Van Geuns*, 988 F.2d 1181, 26 USPQ2d 1057 (Fed. Cir. 1993).

### ***Conclusion***

10. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CAR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CAR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Arun Chakrabarti, Ph.D. whose telephone number is (703) 306-5818.

Art Unit: 1634


If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, W. Gary Jones, can be reached on (703) 308-1152. Any inquiry of a general nature or relating to the status of this application should be directed to the Group analyst Chantae Dessau whose telephone number is (703) 605-1237. Papers related to this application may be submitted to Technology Center 1600 by facsimile transmission via the P.T.O. Fax Center located In Crystal Mall 1. The CM1 Fax Center numbers for Technology Center 1600 are either (703) 305-3014 or (703) 308-4242. Please note that the faxing of such papers must conform with the Notice to Comply published In the Official Gazette, 1096 OG 30 (November 15, 1989).

Arun Chakrabarti

Patent Examiner

Art Unit 1634,

December 17, 2002



W. Gary Jones  
Supervisory Patent Examiner  
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